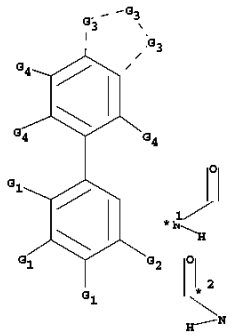
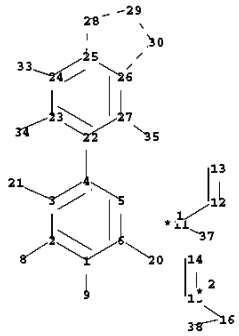




chain nodes :
8 9 11 12 13 14 15 16 20 21 33 34 35 37 38
ring nodes :
1 2 3 4 5 6 22 23 24 25 26 27 28 29 30
chain bonds :
1-9 2-8 3-21 4-22 6-20 11-12 11-37 12-13 14-15 15-16 16-38 23-34 24-33 27-35

ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 22-23 22-27 23-24 24-25 25-26 25-28 26-27 26-30
28-29 29-30
exact/norm bonds :
1-9 2-8 3-21 4-22 6-20 11-12 11-37 12-13 14-15 15-16 16-38 23-34 24-33 25-28
26-30 27-35 28-29 29-30
normalized bonds :
1-2 1-6 2-3 3-4 4-5 5-6 22-23 22-27 23-24 24-25 25-26 26-27

G1:CH3,Et,X,H

G2:[*1],[*2]

G3:C,O,S,N

G4:H,CH3

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 8:CLASS 9:CLASS 11:CLASS 12:CLASS
13:CLASS 14:CLASS 15:CLASS 16:CLASS 20:CLASS 21:CLASS 22:Atom 23:CLASS 24:Atom
25:Atom 26:Atom 27:Atom 28:Atom 29:Atom 30:Atom 33:CLASS 34:CLASS 35:CLASS
37:CLASS 38:CLASS

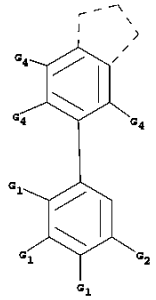
L1 STRUCTURE UPLOADED

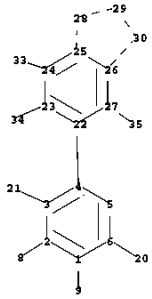
=> d
L1 HAS NO ANSWERS
L1 STR

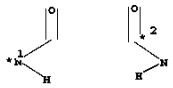
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

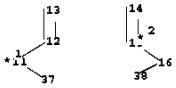
Structure attributes must be viewed using STN Express query preparation.

=>
Uploading C:\Program Files\Stnexp\Queries\10 series\10522955\ALLC123.str









chain nodes :
8 9 11 12 13 14 15 16 20 21 33 34 35 37 38
ring nodes :
1 2 3 4 5 6 22 23 24 25 26 27 28 29 30
chain bonds :
1-9 2-8 3-21 4-22 6-20 11-12 11-37 12-13 14-15 15-16 16-38 23-34 24-33 27-35

ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 22-23 22-27 23-24 24-25 25-26 25-28 26-27 26-30
28-29 29-30
exact/norm bonds :
1-9 2-8 3-21 6-20 11-12 12-13 14-15 15-16 23-34 24-33 25-28 26-30 27-35
28-29 29-30
exact bonds :
4-22 11-37 16-38
normalized bonds :

10/522,955 - Compound Search

1-2 1-6 2-3 3-4 4-5 5-6 22-23 22-27 23-24 24-25 25-26 26-27

G1:CH3,Et,X,H

G2:[*1],[*2]

G3:C,O,S,N

G4:H,CH3

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 8:CLASS 9:CLASS 11:CLASS 12:CLASS
13:CLASS 14:CLASS 15:CLASS 16:CLASS 20:CLASS 21:CLASS 22:Atom 23:CLASS 24:Atom
25:Atom 26:Atom 27:Atom 28:Atom 29:Atom 30:Atom 33:CLASS 34:CLASS 35:CLASS
37:CLASS 38:CLASS

L2   STRUCTURE UPLOADED

=> s l1 not l2
SAMPLE SEARCH INITIATED 10:41:24 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED -   5079 TO ITERATE

 39.4% PROCESSED   2000 ITERATIONS   11 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS: 97307 TO   105853
PROJECTED ANSWERS: 241 TO   875

L3   11 SEA SSS SAM L1 NOT L2

=> s l1 ful
FULL SEARCH INITIATED 10:41:29 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -   102720 TO ITERATE

100.0% PROCESSED   102720 ITERATIONS   540 ANSWERS
SEARCH TIME: 00.00.02

L4   540 SEA SSS FUL L1

=> s l2 ful
FULL SEARCH INITIATED 10:41:35 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -   106172 TO ITERATE

100.0% PROCESSED   106172 ITERATIONS   41 ANSWERS
SEARCH TIME: 00.00.02

L5   41 SEA SSS FUL L2

=> s l4 not l5
L6   499 L4 NOT L5

=> file caplus
COST IN U.S. DOLLARS   SINCE FILE   TOTAL
                                                   ENTRY   SESSION

20 FEB 2008

FULL ESTIMATED COST 356.72 356.93

FILE 'CAPLUS' ENTERED AT 10:41:50 ON 20 FEB 2008
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

FILE COVERS 1907 - 20 Feb 2008 VOL 148 ISS 8
FILE LAST UPDATED: 19 Feb 2008 (20080219/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply.
They are available for your review at:

http://www.cas.org/infopolicy.html

=> s l6
L7 59 L6

=> d scan

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=>
<------User Break------>

=> fil reg
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                         ENTRY    SESSION
FULL ESTIMATED COST                                       0.48     357.41

FILE 'REGISTRY' ENTERED AT 10:42:31 ON 20 FEB 2008
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
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STRUCTURE FILE UPDATES:   19 FEB 2008  HIGHEST RN 1004621-14-0
DICTIONARY FILE UPDATES:  19 FEB 2008  HIGHEST RN 1004621-14-0

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 9, 2008.

  Please note that search-term pricing does apply when
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predicted properties as well as tags indicating availability of
experimental property data in the original document.  For information
on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=> d his

     (FILE 'HOME' ENTERED AT 10:40:11 ON 20 FEB 2008)

     FILE 'REGISTRY' ENTERED AT 10:40:30 ON 20 FEB 2008
L1             STRUCTURE UPLOADED
L2             STRUCTURE UPLOADED
L3          11 S L1 NOT L2
L4         540 S L1 FUL
L5          41 S L2 FUL
L6         499 S L4 NOT L5

     FILE 'CAPLUS' ENTERED AT 10:41:50 ON 20 FEB 2008
L7          59 S L6

     FILE 'REGISTRY' ENTERED AT 10:42:31 ON 20 FEB 2008

=> s l2 full sub=l1
L1 MAY NOT BE USED HERE
The L-number must have been created by a search in this file.  To see
all L-numbers defined in this session, enter DISPLAY HISTORY at an
arrow prompt (=>).  For additional information on subset searching in
this file, enter HELP SUBSET.
ENTER SUBSET L# OR (END):end
SEARCH ENDED BY USER

=> s l2 full sub=l4
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L7 ANSWER 1 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2008:71691 CAPLUS
DOCUMENT NUMBER: 148:168468
TITLE: Trifluoromethyl-substituted benzamides as Eph receptor modulators, their preparation, pharmaceutical compositions, and use in the treatment of neurological disorders
INVENTOR(S): Sivasankaran, Rajeev; Zimmermann, Kaspar
PATENT ASSIGNEE(S): Novartis AG, Switz.; Novartis Pharma GmbH
SOURCE: PCT Int. Appl., 64pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008008821	A2	20080117	WO 2007-US73238	20070711

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.: US 2006-807210P P 20060713

L7 ANSWER 2 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2008:43636 CAPLUS
DOCUMENT NUMBER: 148:121398
TITLE: Cycloalkylcarboxamides and related compounds as modulators of ATP-binding cassette transporters and their preparation, pharmaceutical compositions and use in the treatment of diseases
INVENTOR(S): Hadida Ruah, Sara S.; Miller, Mark T.; Bear, Brian; McCartney, Jason; Grootenhuis, Peter D. J.
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 422pp., Cont.-in-part of U.S. Ser. No. 647,092.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
US 2008009524	A1	20080110	US 2007-824606		20070629
PRIORITY APPLN. INFO.:			US 2005-754558P	P	20051228
			US 2006-802580P	P	20060522
			US 2006-647092	A2	20061228

L7 ANSWER 3 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2008:43435 CAPLUS
DOCUMENT NUMBER: 148:144656
TITLE: Preparation of pyridinonyl PDK1 inhibitors
INVENTOR(S): Lind, Kenneth Egnard; Cao, Kathy; Lin, Edward Yin-Shiang; Nguyen, Thinh Ba; Tangonan, Bradley T.; Erlanson, Daniel A.; Guckian, Kevin; Simmons, Robert Lowell; Lee, Wen-Cherng; Sun, Lihong; Hansen, Stig; Pathan, Nuzhat; Zhang, Lei
PATENT ASSIGNEE(S): Sunesis Pharmaceuticals, USA; Biogen Idec, Inc.
SOURCE: PCT Int. Appl., 311pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008005457	A2	20080110	WO 2007-US15397	20070702

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.:	US 2006-806414P	P	20060630
	US 2007-919057P	P	20070319

L7 ANSWER 4 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:1270696 CAPLUS
DOCUMENT NUMBER: 147:522235
TITLE: Preparation of benzimidazoles as capsaicin receptor VR1 modulators for the treatment of pain
INVENTOR(S): Player, Mark R.; Dax, Scott L.; Parsons, William H.; Brandt, Michael Richard; Calvo, Raul R.; Patel, Sharmila; Liu, Jian; Cheung, Wing S.; Jetter, Michele C.; Lee, Yu-Kai; Youngman, Mark A.; Pan, Wenxi; Weils, Kenneth M.; Beauchamp, Derek A.
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 230pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2007259936	A1	20071108	US 2007-734984	20070413
WO 2007130780	A2	20071115	WO 2007-US66748	20070417

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.: US 2006-797504P P 20060503
OTHER SOURCE(S): MARPAT 147:522235

L7 ANSWER 5 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:1146647 CAPLUS
DOCUMENT NUMBER: 147:448636
TITLE: Preparation of indoles, indazoles, benzimidazoles and their analogs as chemokine receptor CXCR4 and CCR7 inhibitors
INVENTOR(S): Thomas, William D.; Leleti, Manmohan Reddy; Pennell, Andrew M. K.
PATENT ASSIGNEE(S): Chemocentryx, Inc., USA
SOURCE: PCT Int. Appl., 142pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007115231	A2	20071011	WO 2007-US65729	20070330
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RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
US 2007275965	A1	20071129	US 2007-731695	20070330
PRIORITY APPLN. INFO.:			US 2006-787925P P	20060330
OTHER SOURCE(S):	MARPAT 147:448636			

L7 ANSWER 6 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:912148 CAPLUS
DOCUMENT NUMBER: 147:277628
TITLE: Pyrimidinyl benzothiophene compounds as IKKβ kinase inhibitors, their preparation, pharmaceutical compositions, and use in therapy
INVENTOR(S): Dahnke, Karl Robert; Lin, Ho-Shen; Shih, Chuan; Wang, Q. May; Zhang, Bo; Richett, Michael Enrico
PATENT ASSIGNEE(S): Eli Lilly and Company, USA
SOURCE: PCT Int. Appl., 100pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007092095	A2	20070816	WO 2006-US60911	20061115
WO 2007092095	A3	20071108		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRIORITY APPLN. INFO.: US 2005-738097P P 20051118
OTHER SOURCE(S): MARPAT 147:277628

L7 ANSWER 7 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:906796 CAPLUS
DOCUMENT NUMBER: 147:235175
TITLE: Preparation of benzimidazoles, benzoxazoles, benzothiazoles, indoles and their analogs for the treatment of muscular dystrophy and cachexia
INVENTOR(S): Wynne, Graham Michael; Wren, Stephen Paul; Johnson, Peter David; Price, Damien; De Moor, Olivier; Nugent, Gary; Tinsley, Jonathan Mark; Storer, Richard; Mulvaney, Andrew; Pye, Richard Joseph; Dorgan, Colin Richard
PATENT ASSIGNEE(S): Vastox PLC, UK
SOURCE: PCT Int. Appl., 170pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007091106	A2	20070816	WO 2007-GB50055	20070209

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.:	GB 2006-2768	A 20060210
	GB 2006-14690	A 20060724
	GB 2006-19281	A 20060929
	GB 2006-23983	A 20061130

OTHER SOURCE(S): MARPAT 147:235175

L7 ANSWER 8 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:874387 CAPLUS
DOCUMENT NUMBER: 147:257764
TITLE: Preparation of indazole derivatives for treatment of Alzheimer's disease
INVENTOR(S): Churcher, Ian; Choudhury, Hedaythul; Hunt, Peter; Jelley, Richard; Nadin, Alan; Nanthakumar, Carmel B.; Simpson, Peter Brian; Wilkie, Neil
PATENT ASSIGNEE(S): Merck Sharp & Dohme Limited, UK
SOURCE: PCT Int. Appl., 59pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007088401	A1	20070809	WO 2007-GB50048	20070202
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				

PRIORITY APPLN. INFO.: GB 2006-2178 A 20060203
OTHER SOURCE(S): MARPAT 147:257764
REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 9 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:846121 CAPLUS
DOCUMENT NUMBER: 147:211534
TITLE: Cycloalkylcarboxamides and related compounds as modulators of ATP-binding cassette transporters and their preparation, pharmaceutical compositions and use in the treatment of diseases
INVENTOR(S): Ruah, Sara S. Hadida; Miller, Mark T.; Bear, Brian; McCartney, Jason; Grootenhuis, Peter D. J.
PATENT ASSIGNEE(S): Vertex Pharmaceuticals Incorporated, USA
SOURCE: PCT Int. Appl., 249pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007087066	A2	20070802	WO 2006-US49412	20061228
WO 2007087066	A3	20071025		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRIORITY APPLN. INFO.: US 2005-754558P P 20051228
US 2006-802580P P 20060522
OTHER SOURCE(S): MARPAT 147:211534

L7 ANSWER 10 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:670446 CAPLUS
DOCUMENT NUMBER: 147:95910
TITLE: Preparation of proline amides for treating Flaviviridae family virus infection
INVENTOR(S): Schmitz, Franz Ulrich; Roberts, Christopher Don; Abadi, Ali Dehghani Mohammad; Griffith, Ronald Conrad; Leivers, Martin Robert
PATENT ASSIGNEE(S): Genelabs Technologies, Inc., USA
SOURCE: PCT Int. Appl., 115pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007070556	A2	20070621	WO 2006-US47503	20061212
WO 2007070556	A3	20070830		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA			
US 2007265262	A1	20071115	US 2006-609854	20061212
PRIORITY APPLN. INFO.:			US 2005-749771P P	20051212
OTHER SOURCE(S):	MARPAT 147:95910			

L7 ANSWER 11 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:510466 CAPLUS
DOCUMENT NUMBER: 146:501048
TITLE: Preparation of heterocyclic amide compounds as FXR inhibitors
INVENTOR(S): Miura, Shotaro; Shimada, Mitsuyuki; Marui, Shogo; Tamura, Norikazu; Nakada, Yoshihisa; Tozawa, Ryuichi; Sakamoto, Junichi; Funabashi, Yasunori; Hosono, Hiroshi
PATENT ASSIGNEE(S): Takeda Pharmaceutical Company Limited, Japan
SOURCE: PCT Int. Appl., 1320pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007052843	A1	20070510	WO 2006-JP322420	20061102

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.: JP 2005-321600 A 20051104
JP 2006-251883 A 20060915
OTHER SOURCE(S): MARPAT 146:501048
REFERENCE COUNT: 83 THERE ARE 83 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 12 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:384878 CAPLUS
DOCUMENT NUMBER: 146:411416
TITLE: Silver halide color photosensitive material containing fluorine compound
INVENTOR(S): Yokota, Koichi
PATENT ASSIGNEE(S): Fuji Photo Film Co., Ltd., Japan
SOURCE: U.S. Pat. Appl. Publ., 53pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2007077527	A1	20070405	US 2006-529522	20060929
US 7306900	B2	20071211		
JP 2007094243	A	20070412	JP 2005-286065	20050930
PRIORITY APPLN. INFO.:			JP 2005-286065 A	20050930

OTHER SOURCE(S): MARPAT 146:411416

L7 ANSWER 13 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:14480 CAPLUS
DOCUMENT NUMBER: 146:121821
TITLE: Preparation of bicyclic derivatives as p38 kinase inhibitors
INVENTOR(S): Almansa Rosales, Carmen; Virgili Bernado, Marina
PATENT ASSIGNEE(S): J. Uriach y Compania S.A., Spain
SOURCE: PCT Int. Appl., 80pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007000339	A1	20070104	WO 2006-EP6255	20060628
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2006263961	A1	20070104	AU 2006-263961	20060628
PRIORITY APPLN. INFO.:			EP 2005-380140 A	20050629
			WO 2006-EP6255 W	20060628

OTHER SOURCE(S): MARPAT 146:121821
REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 14 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:1252528 CAPLUS
DOCUMENT NUMBER: 146:27971
TITLE: Preparation of 6,7-unsaturated-7-carbamoyl substituted morphinan derivatives as opioid antagonists for the treatment of nausea, vomiting and/or constipation
INVENTOR(S): Inagaki, Masanao; Hara, Shin-Ichiro; Haga, Nobuhiro; Tamura, Yoshinori; Goto, Yoshihisa; Hasegawa, Tsuyoshi
PATENT ASSIGNEE(S): Shionogi & Co., Ltd., Japan
SOURCE: PCT Int. Appl., 162pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006126637	A1	20061130	WO 2006-JP310454	20060525
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
WO 2006126529	A1	20061130	WO 2006-JP310231	20060523
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2006250390	A1	20061130	AU 2006-250390	20060525
PRIORITY APPLN. INFO.:			JP 2005-151864 A	20050525
			JP 2006-65762 A	20060310
			WO 2006-JP310231 A	20060523
			WO 2006-JP310454 W	20060525

OTHER SOURCE(S): MARPAT 146:27971
REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 15 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:1251694 CAPLUS
DOCUMENT NUMBER: 146:27969
TITLE: Preparation of 6,7-unsaturated-7-carbamoyl substituted morphinan derivatives as opioid antagonists for the treatment of nausea, vomiting and/or constipation
INVENTOR(S): Inagaki, Masanao; Hara, Shin-Ichiro; Haga, Nobuhiro; Tamura, Yoshinori; Goto, Yoshihisa; Hasegawa, Tsuyoshi
PATENT ASSIGNEE(S): Shionogi & Co., Ltd., Japan
SOURCE: PCT Int. Appl., 145pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006126529	A1	20061130	WO 2006-JP310231	20060523
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2006250390	A1	20061130	AU 2006-250390	20060525
WO 2006126637	A1	20061130	WO 2006-JP310454	20060525
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				

PRIORITY APPLN. INFO.:		
JP 2005-151864	A	20050525
JP 2006-65762	A	20060310
WO 2006-JP310231	W	20060523
WO 2006-JP310454	W	20060525

OTHER SOURCE(S): MARPAT 146:27969
REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 16 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:1226132 CAPLUS
DOCUMENT NUMBER: 146:7967
TITLE: Preparation of benzo[d]isoxazol-3-ylamines as vanilloid receptor 1 inhibitors
INVENTOR(S): Frank, Robert; Merla, Beatrix; Reich, Melanie; Jostock, Ruth
PATENT ASSIGNEE(S): Gruenenthal GmbH, Germany
SOURCE: PCT Int. Appl., 113pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006122799	A1	20061123	WO 2006-EP4698	20060518
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
DE 102005038947	A1	20061130	DE 2005-102005038947	20050816
CA 2608386	A1	20061123	CA 2006-2608386	20060518
PRIORITY APPLN. INFO.:			DE 2005-102005023589A	20050518
			DE 2005-102005038947A	20050816
			WO 2006-EP4698 W	20060518

OTHER SOURCE(S): MARPAT 146:7967
REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 17 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:888369 CAPLUS
DOCUMENT NUMBER: 145:293091
TITLE: Preparation of bicyclic heteroaromatic derivatives as anticancer agents
INVENTOR(S): Kauffman, Goss Stryker; Li, Chao; Lippa, Blaise Scott; Morris, Joel; Pan, Gonghua
PATENT ASSIGNEE(S): Pfizer Products Inc., USA
SOURCE: PCT Int. Appl., 152pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006090261	A1	20060831	WO 2006-IB406	20060215
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
CA 2598956	A1	20060831	CA 2006-2598956	20060215
EP 1858902	A1	20071128	EP 2006-710461	20060215
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				

PRIORITY APPLN. INFO.: US 2005-656467P P 20050224
WO 2006-IB406 W 20060215
OTHER SOURCE(S): MARPAT 145:293091
REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 18 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:845716 CAPLUS
DOCUMENT NUMBER: 145:293345
TITLE: Preparation of N-acyl-amino acid derivatives for controlling function of GPR34 receptor as antagonists or inverse agonists
INVENTOR(S): Ito, Fumio; Kimura, Eiji; Imai, Tomomi; Mori, Masaaki; Aramaki, Yoshio; Kohara, Yasuhisa; Sugo, Tsukasa; Hayase, Yoji; Kobayashi, Hiromi; Ogi, Kazuhiro
PATENT ASSIGNEE(S): Takeda Pharmaceutical Company Limited, Japan
SOURCE: PCT Int. Appl., 597pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006088246	A1	20060824	WO 2006-JP303357	20060217

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

EP 1849465	A1	20071031	EP 2006-714496	20060217

R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR

PRIORITY APPLN. INFO.:		
JP 2005-41775	A	20050218
JP 2005-315146	A	20051028
WO 2006-JP303357	W	20060217

OTHER SOURCE(S): MARPAT 145:293345
REFERENCE COUNT: 28 THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 19 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:844783 CAPLUS
DOCUMENT NUMBER: 145:271746
TITLE: Fused heterocyclic compounds useful as inhibitors of histone deacetylase
INVENTOR(S): Anandan, Sampath K.; Xiao, Xiao-Yi; Ward, John S.; Patel, Dinesh V.
PATENT ASSIGNEE(S): Mikana Therapeutics, Inc., USA
SOURCE: PCT Int. Appl., 223pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006088949	A1	20060824	WO 2006-US5312	20060214
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2006214319	A1	20060824	AU 2006-214319	20060214
CA 2596015	A1	20060824	CA 2006-2596015	20060214
US 2006199829	A1	20060907	US 2006-354594	20060214
EP 1851219	A1	20071107	EP 2006-735122	20060214
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
PRIORITY APPLN. INFO.:			US 2005-652870P	P 20050214
			WO 2006-US5312	W 20060214

OTHER SOURCE(S): MARPAT 145:271746
REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 20 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:606657 CAPLUS
DOCUMENT NUMBER: 145:83212
TITLE: Preparation of indolinones as antiproliferative agents
INVENTOR(S): McConnell, Darryl; Weyer-Czernilofsky, Ulrike; Impagnatiello, Maria; Steurer, Steffen; Brueckner, Ralph; Krist, Bernd; Betzemeier, Bodo; Hilberg, Frank; Heckel, Armin; Roth, Gerald Juergen; Kley, Joerg; Lehmann-Lintz, Thorsten
PATENT ASSIGNEE(S): Boehringer Ingelheim International G.m.b.H., Germany; Boehringer Ingelheim Pharma Gmbh & Co. K.-G.
SOURCE: PCT Int. Appl., 90 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006064044	A1	20060622	WO 2005-EP56821	20051215
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
CA 2589501	A1	20060622	CA 2005-2589501	20051215
EP 1828123	A1	20070905	EP 2005-817469	20051215
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
US 2006135592	A1	20060622	US 2005-303168	20051216
US 2007088051	A1	20070419	US 2006-558953	20061113
PRIORITY APPLN. INFO.:			EP 2004-29981	A 20041217
			WO 2005-EP56821	W 20051215
			US 2005-303168	B1 20051216

OTHER SOURCE(S): MARPAT 145:83212
REFERENCE COUNT: 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 21 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:513602 CAPLUS
DOCUMENT NUMBER: 145:46271
TITLE: Preparation of glycopeptide antibiotic monomer derivatives having antibacterial activity against vancomycin-resistant bacteria
INVENTOR(S): Arimoto, Hirokazu; Lu, Jun; Yamano, Yoshinori; Yasukata, Tatsuro; Yoshida, Osamu; Iwaki, Tsutomu; Yoshida, Yutaka; Kato, Issei; Morimoto, Kenji; Yasoshima, Kayo
PATENT ASSIGNEE(S): National University Corporation Nagoya University, Japan; Shionogi & Co., Ltd.
SOURCE: PCT Int. Appl., 244 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006057303	A1	20060601	WO 2005-JP21587	20051124
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2005308160	A1	20060601	AU 2005-308160	20051124
CA 2588285	A1	20060601	CA 2005-2588285	20051124
EP 1818340	A1	20070815	EP 2005-809139	20051124
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
IN 2007CN02297	A	20070907	IN 2007-CN2297	20070529
KR 2007092719	A	20070913	KR 2007-714842	20070628
CN 101111513	A	20080123	CN 2005-80047421	20070730

PRIORITY APPLN. INFO.: JP 2004-344231 A 20041129
JP 2005-212471 A 20050722
WO 2005-JP21587 W 20051124

OTHER SOURCE(S): MARPAT 145:46271
REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 22 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:469591 CAPLUS
DOCUMENT NUMBER: 144:488517
TITLE: Preparation of 6-substituted 2-oxo-3-(1H-pyrrol-2-ylmethylene)-2,3-dihydro-1H-indole derivatives and their compositions as protein kinase inhibitors
INVENTOR(S): Wan, Yongqin; Mi, Yuan; Fan, Yi; Cheng, Dai; Liu, Yi; Gray, Nathanael Schiander; Albaugh, Pamela A.
PATENT ASSIGNEE(S): Irm LLC, Bermuda
SOURCE: PCT Int. Appl., 108 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006052936	A2	20060518	WO 2005-US40372	20051107
WO 2006052936	A3	20061026		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2005304719	A1	20060518	AU 2005-304719	20051107
CA 2583737	A1	20060518	CA 2005-2583737	20051107
EP 1814545	A2	20070808	EP 2005-851419	20051107
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
CN 101056632	A	20071017	CN 2005-80038332	20051107
IN 2007DN03051	A	20070831	IN 2007-DN3051	20070424
KR 2007084066	A	20070824	KR 2007-710439	20070508
NO 2007002887	A	20070803	NO 2007-2887	20070606

PRIORITY APPLN. INFO.:	US 2004-626785P	P	20041109
	US 2005-709648P	P	20050819
	WO 2005-US40372	W	20051107

OTHER SOURCE(S): MARPAT 144:488517

L7 ANSWER 23 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:437473 CAPLUS
DOCUMENT NUMBER: 144:425684
TITLE: Imidazole derivatives
INVENTOR(S): Campbell, David; Betancort, Juan
PATENT ASSIGNEE(S): Phenomix Corporation, USA
SOURCE: PCT Int. Appl., 47 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006050162	A2	20060511	WO 2005-US39074	20051028
WO 2006050162	A3	20070201		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
US 2006135553	A1	20060622	US 2005-262394	20051028
PRIORITY APPLN. INFO.:			US 2004-623421P P	20041028
OTHER SOURCE(S):	MARPAT 144:425684			

L7 ANSWER 24 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:149229 CAPLUS
DOCUMENT NUMBER: 144:233084
TITLE: Trifluoromethyl-substituted benzamides as kinase inhibitors and their preparation, pharmaceutical compositions, and use for treatment of proliferative diseases
INVENTOR(S): Caravatti, Giorgio; Furet, Pascal; Imbach, Patricia; Martiny-Baron, Georg; Perez, Lawrence Blas; Sheng, Tao
PATENT ASSIGNEE(S): Novartis AG, Switz.; Novartis Pharma GmbH
SOURCE: PCT Int. Appl., 81 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006015859	A1	20060216	WO 2005-EP8695	20050810
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2005270313	A1	20060216	AU 2005-270313	20050810
CA 2575316	A1	20060216	CA 2005-2575316	20050810
US 2006035897	A1	20060216	US 2005-201348	20050810
EP 1778640	A1	20070502	EP 2005-777531	20050810
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, HR				
CN 101039914	A	20070919	CN 2005-80034662	20050810
IN 2007DN00848	A	20070803	IN 2007-DN848	20070131
KR 2007046851	A	20070503	KR 2007-703238	20070209
NO 2007001300	A	20070419	NO 2007-1300	20070309

PRIORITY APPLN. INFO.: GB 2004-17905 A 20040811
WO 2005-EP8695 W 20050810
OTHER SOURCE(S): MARPAT 144:233084
REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 25 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:75243 CAPLUS
DOCUMENT NUMBER: 144:150384
TITLE: Preparation of 1,4-disubstituted naphthalenes as inhibitors of p38 MAP kinase
INVENTOR(S): Ashwell, Mark Antony; Liu, Yanbin; Ali, Syed; Hill, Jason; Wrona, Woj
PATENT ASSIGNEE(S): Arqule, Inc., USA
SOURCE: PCT Int. Appl., 261 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006010082	A1	20060126	WO 2005-US24441	20050708
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
US 2008032967	A1	20080207	US 2007-631617	20070720
PRIORITY APPLN. INFO.:			US 2004-585862P P	20040708
			WO 2005-US24441 W	20050708

OTHER SOURCE(S): MARPAT 144:150384
REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 26 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:1350605 CAPLUS
DOCUMENT NUMBER: 144:69837
TITLE: Preparation of 3-aminoindazoles as serum and glucocorticoid-regulated kinase (SGK) inhibitors
INVENTOR(S): Dorsch, Dieter; Burgdorf, Lars Thore; Gericke, Rolf; Beier, Norbert; Mederski, Werner; Lang, Florian
PATENT ASSIGNEE(S): Merck Patent GmbH, Germany
SOURCE: PCT Int. Appl., 136 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005123688	A2	20051229	WO 2005-EP3513	20050404
WO 2005123688	A3	20060223		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
DE 102004028862	A1	20051229	DE 2004-102004028862	20040615
AU 2005254617	A1	20051229	AU 2005-254617	20050404
CA 2570264	A1	20051229	CA 2005-2570264	20050404
EP 1765788	A2	20070328	EP 2005-729376	20050404
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, LV				
JP 2008502610	T	20080131	JP 2007-515792	20050404
IN 2006KN03519	A	20070615	IN 2006-KN3519	20061124
US 2007232620	A1	20071004	US 2006-629504	20061214
PRIORITY APPLN. INFO.:			DE 2004-102004028862A	20040615
			WO 2005-EP3513 W	20050404

OTHER SOURCE(S): MARPAT 144:69837

L7 ANSWER 27 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:1154550 CAPLUS
DOCUMENT NUMBER: 143:422508
TITLE: Preparation of 2-(1-azabicyclo[2.2.2]oct-3-yl)-2,3-dihydroisoindol-1-one and 5-(1-azabicyclo[2.2.2]oct-3-yl)-5,6-dihydro-furo[2,3-c]pyrrol-4-one derivatives for therapeutic use as ligands for the α7 nicotinic acetylcholine receptor (α7nAChR)
INVENTOR(S): Chapdelaine, Marc; Herzog, Keith J.
PATENT ASSIGNEE(S): Astrazeneca AB, Swed.; Chapdelaine, Marc; Herzog, Keith J.
SOURCE: PCT Int. Appl., 59 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005100351	A1	20051027	WO 2005-SE500	20050406
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2005233492	A1	20051027	AU 2005-233492	20050406
CA 2563010	A1	20051027	CA 2005-2563010	20050406
EP 1737854	A1	20070103	EP 2005-722314	20050406
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, LV, MK, YU				
CN 1968951	A	20070523	CN 2005-80019493	20050406
BR 2005009777	A	20071023	BR 2005-9777	20050406
JP 2007532637	T	20071115	JP 2007-508300	20050406
IN 2006DN05559	A	20070831	IN 2006-DN5559	20060925
MX 2006PA11725	A	20061211	MX 2006-PA11725	20061010
US 2007213342	A1	20070913	US 2006-599839	20061011
KR 2007020445	A	20070221	KR 2006-721260	20061013
NO 2006005199	A	20061113	NO 2006-5199	20061113
PRIORITY APPLN. INFO.:			SE 2004-970 A	20040414
			WO 2005-SE500 W	20050406

OTHER SOURCE(S): MARPAT 143:422508
REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 28 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:962210 CAPLUS
DOCUMENT NUMBER: 143:266917
TITLE: Preparation of azolone derivatives for the treatment of anxiety and depression
INVENTOR(S): Kodo, Toru; Fukaya, Takayuki; Koyama, Koji; Masumoto, Shuji; Fujibayashi, Nao
PATENT ASSIGNEE(S): Sumitomo Pharmaceuticals Co., Ltd., Japan
SOURCE: PCT Int. Appl., 161 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005080334	A1	20050901	WO 2005-JP3095	20050218
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2005214258	A1	20050901	AU 2005-214258	20050218
CA 2554774	A1	20050901	CA 2005-2554774	20050218
EP 1719761	A1	20061108	EP 2005-710695	20050218
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS				
CN 1922141	A	20070228	CN 2005-80005614	20050218
US 2007191447	A1	20070816	US 2006-590157	20060821
IN 2006CN03053	A	20070608	IN 2006-CN3053	20060822
PRIORITY APPLN. INFO.:			JP 2004-45979 A	20040223
			WO 2005-JP3095 W	20050218

OTHER SOURCE(S): MARPAT 143:266917
REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 29 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:732643 CAPLUS
DOCUMENT NUMBER: 143:193999
TITLE: Preparation of fused heteroaryl derivatives as p38 kinase inhibitors
INVENTOR(S): Campos, Sebastien Andre; Swanson, Stephen; Walker, Ann Louise
PATENT ASSIGNEE(S): Smithkline Beecham Corporation, USA
SOURCE: PCT Int. Appl., 59 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005073219	A1	20050811	WO 2005-GB281	20050127
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1745038	A1	20070124	EP 2005-702034	20050127
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, HR, LV				
JP 2007519695	T	20070719	JP 2006-550298	20050127
US 2007142372	A1	20070621	US 2006-587614	20060728
PRIORITY APPLN. INFO.:			GB 2004-2140 A	20040130
			WO 2005-GB281 W	20050127

OTHER SOURCE(S): CASREACT 143:193999; MARPAT 143:193999
REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 30 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:729633 CAPLUS
DOCUMENT NUMBER: 143:211906
TITLE: Preparation of fused heteroaryl derivatives as p38 kinase inhibitors
INVENTOR(S): Bamborough, Paul; Campos, Sebastien Andre; Patel, Vipulkumar Kantibhai; Swanson, Stephen; Walker, Ann Louise
PATENT ASSIGNEE(S): Smithkline Beecham Corporation, USA
SOURCE: PCT Int. Appl., 123 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005073189	A1	20050811	WO 2005-GB265	20050127
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1708996	A1	20061011	EP 2005-702022	20050127
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, HR, IS				
JP 2007519692	T	20070719	JP 2006-550294	20050127
PRIORITY APPLN. INFO.:			GB 2004-2143	A 20040130
			WO 2005-GB265	W 20050127

OTHER SOURCE(S): MARPAT 143:211906
REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 31 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:588997 CAPLUS
DOCUMENT NUMBER: 143:115438
TITLE: Preparation of substituted indol-2-ols as kinase inhibitors
INVENTOR(S): Gangloff, Anthony R.; Nowakowski, Jacek; Paraselli, Bheema R.; Stafford, Jeffrey A.; Tennant, Michael G.
PATENT ASSIGNEE(S): Syrrx, Inc., USA
SOURCE: PCT Int. Appl., 179 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005061519	A1	20050707	WO 2004-US42631	20041217
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2005153966	A1	20050714	US 2004-15348	20041217
EP 1694686	A1	20060830	EP 2004-814774	20041217
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS				
JP 2007514759	T	20070607	JP 2006-545517	20041217
PRIORITY APPLN. INFO.:			US 2003-531202P P	20031219
			WO 2004-US42631 W	20041217

OTHER SOURCE(S): MARPAT 143:115438
REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 32 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:141046 CAPLUS
DOCUMENT NUMBER: 142:219141
TITLE: Preparation of benzofuran derivatives useful for treating hyperproliferative disorders
INVENTOR(S): Zhang, Chengzhi; Dumas, Jacques; Ladouceur, Gaetan H.; Zhao, Qian; Hentemann, Martin F.; Verma, Sharad K.; Zhu, Qingming; Lavoie, Rico C.; Fan, Jianmei; Phillips, Barton
PATENT ASSIGNEE(S): Bayer Pharmaceuticals Corporation, USA
SOURCE: PCT Int. Appl., 128 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005014566	A1	20050217	WO 2004-US25480	20040806
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2534678	A1	20050217	CA 2004-2534678	20040806
EP 1654245	A1	20060510	EP 2004-780334	20040806
R: DE, ES, FR, GB, IT				
JP 2007501796	T	20070201	JP 2006-522757	20040806
US 2006194816	A1	20060831	US 2006-566343	20060127
PRIORITY APPLN. INFO.:			US 2003-494165P P	20030807
			WO 2004-US25480 W	20040806

OTHER SOURCE(S): CASREACT 142:219141; MARPAT 142:219141
REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 33 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:55229 CAPLUS
DOCUMENT NUMBER: 142:134457
TITLE: Preparation of substituted indoles as inhibitors of microsomal PGE synthase-1 for the treatment of inflammation
INVENTOR(S): Olofsson, Kristofer; Suna, Edgars; Pelcman, Benjamin; Ozola, Vita; Katkevics, Martins; Kalvins, Ivars
PATENT ASSIGNEE(S): Biolipox AB, Swed.; McNeeney, Stephen Phillip; Schaal, Wesley
SOURCE: PCT Int. Appl., 131 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

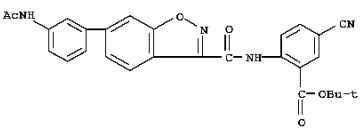
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005005415	A1	20050120	WO 2004-GB2996	20040709
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2528626	A1	20050120	CA 2004-2528626	20040709
EP 1646624	A1	20060419	EP 2004-743337	20040709
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
JP 2007516203	T	20070621	JP 2006-518373	20040709
US 2006160879	A1	20060720	US 2006-563464	20060214
PRIORITY APPLN. INFO.:			SE 2003-2035 A	20030709
			US 2003-485390P P	20030709
			WO 2004-GB2996 W	20040709

OTHER SOURCE(S): CASREACT 142:134457; MARPAT 142:134457
REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 36 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:182843 CAPLUS
DOCUMENT NUMBER: 140:235498
TITLE: Preparation of antibacterial benzoic acid derivatives
INVENTOR(S): Thorarensen, Atli; Ruble, Craig J.; Fisher, Jed F.; Romero, Donna L.; Beauchamp, Thomas J.; Northuis, Jill M.
PATENT ASSIGNEE(S): Pharmacia & Upjohn Company, USA
SOURCE: PCT Int. Appl., 500 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

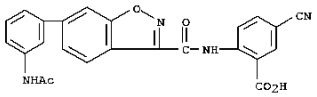
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004018428	A1	20040304	WO 2003-US24796	20030822
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2004110802	A1	20040610	US 2003-645802	20030820
AU 2003264005	A1	20040311	AU 2003-264005	20030822
PRIORITY APPLN. INFO.:			US 2002-405429P P	20020823
			US 2002-430592P P	20021203
			WO 2003-US24796 W	20030822

OTHER SOURCE(S): MARPAT 140:235498
IT 668970-25-0P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of benzoic acid derivs. as antibacterial agents)
RN 668970-25-0 CAPLUS
CN Benzoic acid, 2-[[[6-[3-(acetylamino)phenyl]-1,2-benzisoxazol-3-yl]carbonyl]amino]-5-cyano-, 1,1-dimethylethyl ester (CA INDEX NAME)

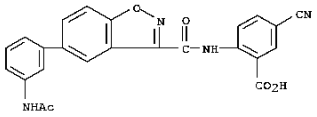


IT 668970-23-8P 668970-28-3P
RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of benzoic acid derivs. as antibacterial agents)

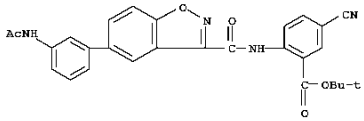
L7 ANSWER 36 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)
RN 668970-23-8 CAPLUS
CN Benzoic acid, 2-[[[6-[3-(acetylamino)phenyl]-1,2-benzisoxazol-3-yl]carbonyl]amino]-5-cyano- (CA INDEX NAME)



RN 668970-28-3 CAPLUS
CN Benzoic acid, 2-[[[5-[3-(acetylamino)phenyl]-1,2-benzisoxazol-3-yl]carbonyl]amino]-5-cyano- (CA INDEX NAME)



IT 668970-29-4
RL: RCT (Reactant); RACT (Reactant or reagent)
(reactant; preparation of benzoic acid derivs. as antibacterial agents)
RN 668970-29-4 CAPLUS
CN Benzoic acid, 2-[[[5-[3-(acetylamino)phenyl]-1,2-benzisoxazol-3-yl]carbonyl]amino]-5-cyano-, 1,1-dimethylethyl ester (CA INDEX NAME)



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

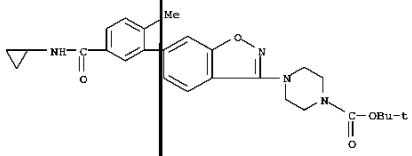
INSTANT

L7 ANSWER 37 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:100989 CAPLUS
DOCUMENT NUMBER: 140:146133
TITLE: Preparation of fused heteroaryls, in particular benzisoxazoles and indazoles, for use as p38 kinase inhibitors in the treatment of rheumatoid arthritis
INVENTOR(S): Angell, Richard Martyn; Baldwin, Ian Robert; Bamborough, Paul; Deboeck, Nigel Marc; Longstaff, Timothy; Swanson, Stephen
PATENT ASSIGNEE(S): Smithkline Beecham Corporation, USA
SOURCE: PCT Int. Appl., 135 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

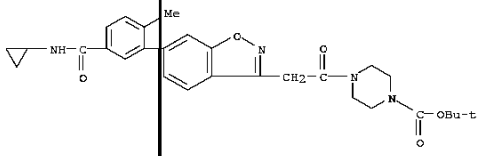
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004010995	A1	20040205	WO 2003-GB3316	20030730
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003248978	A1	20040216	AU 2003-248978	20030730
EP 1531812	A1	20050525	EP 2003-771208	20030730
EP 1531812	B1	20070627		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2005538100	T	20051215	JP 2004-523985	20030730
AT 365551	T	20070715	AT 2003-771208	20030730
ES 2289336	T3	20080201	ES 2003-771208	20030730
US 2006122221	A1	20060608	US 2005-522955	20051114
PRIORITY APPLN. INFO.:			GB 2002-17757 A	20020731
			WO 2003-GB3316 W	20030730

OTHER SOURCE(S): MARPAT 140:146133
IT 651780-05-1P, 1,1-Dimethylethyl 4-[6-[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]-1,2-benzisoxazol-3-yl]-1-piperazinecarboxylate 651780-09-5P 651780-39-1P, N-Cyclopropyl-3-[3-[(4-fluorophenyl)sulfonyl]-1-[[[2-(trimethylsilyl)ethyl]oxy]methyl]-1H-indazol-6-yl]-4-methylbenzamide 651780-45-9P, N-Cyclopropyl-3-fluoro-4-methyl-5-[3-(methylsulfonyl)-1-[[[2-(trimethylsilyl)ethyl]oxy]methyl]-1H-indazol-6-yl]benzamide
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of fused heteroaryls as p38 kinase inhibitors for treatment of rheumatoid arthritis)
RN 651780-05-1 CAPLUS
CN 1-Piperazinecarboxylic acid, 4-[6-[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]-1,2-benzisoxazol-3-yl]-, 1,1-dimethylethyl ester (CA INDEX NAME)

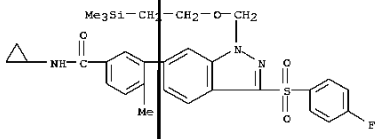
L7 ANSWER 37 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)



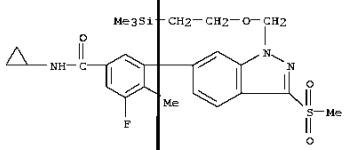
RN 651780-09-5 CAPLUS
CN 1-Piperazinecarboxylic acid, 4-[[6-[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]-1,2-benzisoxazol-3-yl]acetyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RN 651780-39-1 CAPLUS
CN Benzamide, N-cyclopropyl-3-[3-[(4-fluorophenyl)sulfonyl]-1-[[2-(trimethylsilyl)ethoxy]methyl]-1H-indazol-6-yl]-4-methyl- (CA INDEX NAME)



RN 651780-45-9 CAPLUS
CN Benzamide, N-cyclopropyl-3-fluoro-4-methyl-5-[3-(methylsulfonyl)-1-[[2-(trimethylsilyl)ethoxy]methyl]-1H-indazol-6-yl]- (CA INDEX NAME)

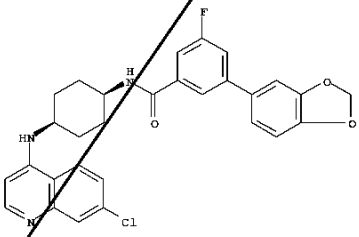


L7 ANSWER 38 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2003:971736 CAPLUS
DOCUMENT NUMBER: 140:16656
TITLE: cis-N-(Quinolin-4-yl)cyclohexane-1,4-diamine derivatives as antagonists of melanin concentrating hormone (MCH) and their pharmaceutical compositions and therapeutic uses, e.g., for treatment of obesity
INVENTOR(S): Kym, Philip R.; Hartandi, Kresna; Gao, Ju; Phelan, Kathleen M.; Akritopoulou-Zanze, Irini; Collins, Christine A.; Vasudevan, Anil; Verzal, Mary K.
PATENT ASSIGNEE(S): Abbott Laboratories, USA
SOURCE: U.S. Pat. Appl. Publ., 89 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003229119	A1	20031211	US 2003-372359	20030221
US 6818772	B2	20041116		
PRIORITY APPLN. INFO.:			US 2002-359081P P	20020222

OTHER SOURCE(S): MARPAT 140:16656
IT 589493-31-2P, cis-3-(1,3-Benzodioxol-5-yl)-N-[4-[(7-chloroquinolin-4-yl)amino]cyclohexyl]-5-fluorobenzamide
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of quinolinylcyclohexanediamine derivs. as MCH receptor antagonists)
RN 589493-31-2 CAPLUS
CN Benzamide, 3-(1,3-benzodioxol-5-yl)-N-[cis-4-[(7-chloro-4-quinolinyl)amino]cyclohexyl]-5-fluoro- (CA INDEX NAME)

Relative stereochemistry.



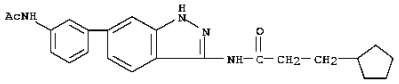
REFERENCE COUNT: 14 THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 39 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2003:931339 CAPLUS
DOCUMENT NUMBER: 140:5044
TITLE: Preparation of 3-aminoindazole derivatives as kinase inhibitors
INVENTOR(S): Martina, Katia; Brill, Wolfgang
PATENT ASSIGNEE(S): Pharmacia Italia S.P.A., Italy
SOURCE: PCT Int. Appl., 99 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

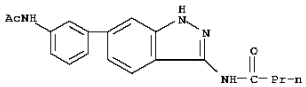
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003097610	A1	20031127	WO 2003-EP4862	20030508
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2486101	A1	20031127	CA 2003-2486101	20030508
AU 2003227741	A1	20031202	AU 2003-227741	20030508
EP 1506176	A1	20050216	EP 2003-725180	20030508
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003011291	A	20050329	BR 2003-11291	20030508
JP 2005534635	T	20051117	JP 2004-505343	20030508
MX 2004PA11417	A	20050214	MX 2004-PA11417	20041117
US 2006106083	A1	20060518	US 2004-990866	20041117
PRIORITY APPLN. INFO.:			US 2002-381092P P	20020517
			WO 2003-EP4862 W	20030508

OTHER SOURCE(S): CASREACT 140:5044; MARPAT 140:5044
IT 627848-14-0P 627848-15-1P 627848-16-2P
627848-91-3P 627848-92-4P 627848-93-5P
627849-36-9P 627849-37-0P 627858-25-7P
627858-26-8P 627858-27-9P 627858-28-0P
627858-29-1P 627858-30-4P 627858-31-5P
627858-32-6P 627858-33-7P 627858-34-8P
627859-06-7P 627859-07-8P 627859-08-9P
627859-09-0P 627859-10-3P 627859-11-4P
627859-12-5P 627859-13-6P 627859-14-7P
RL: CPN (Combinatorial preparation); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); CMBI (Combinatorial study); PREP (Preparation); USES (Uses)
(preparation of 3-aminoindazole derivs. as kinase inhibitors)
RN 627848-14-0 CAPLUS
CN Cyclopentanepropanamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]- (CA INDEX NAME)

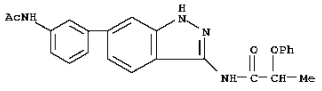
L7 ANSWER 39 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)



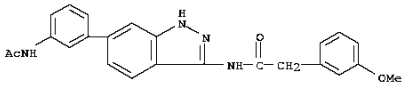
RN 627848-15-1 CAPLUS
CN Butanamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]- (CA INDEX NAME)



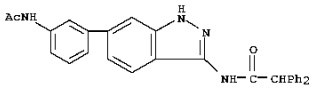
RN 627848-16-2 CAPLUS
CN Propanamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]-2-phenoxy- (CA INDEX NAME)



RN 627848-91-3 CAPLUS
CN Benzeneacetamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]-3-methoxy- (CA INDEX NAME)

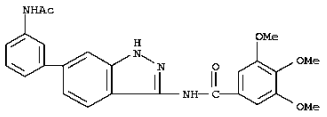


RN 627848-92-4 CAPLUS
CN Benzeneacetamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]-α-phenyl- (CA INDEX NAME)

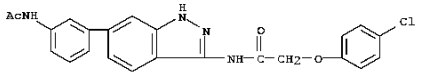


RN 627848-93-5 CAPLUS

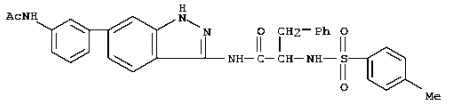
L7 ANSWER 39 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)
CN Benzamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]-3,4,5-trimethoxy- (CA INDEX NAME)



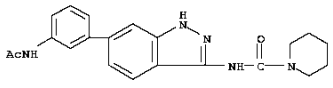
RN 627849-36-9 CAPLUS
CN Acetamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]-2-(4-chlorophenoxy)- (CA INDEX NAME)



RN 627849-37-0 CAPLUS
CN Benzenepropanamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]-α-[[(4-methylphenyl)sulfonyl]amino]- (CA INDEX NAME)

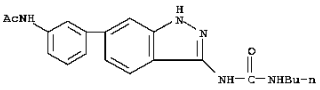


RN 627858-25-7 CAPLUS
CN 1-Piperidinecarboxamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]- (CA INDEX NAME)

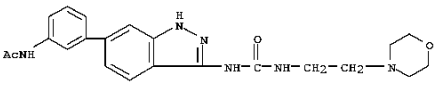


RN 627858-26-8 CAPLUS
CN Acetamide, N-[3-[3-[[(butylamino)carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

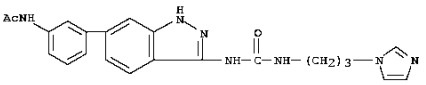
L7 ANSWER 39 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)



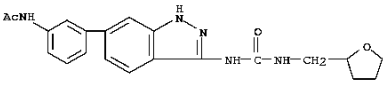
RN 627858-27-9 CAPLUS
CN Acetamide, N-[3-[3-[[[[2-(4-morpholinyl)ethyl]amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)



RN 627858-28-0 CAPLUS
CN Acetamide, N-[3-[3-[[[[3-(1H-imidazol-1-yl)propyl]amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

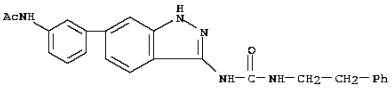


RN 627858-29-1 CAPLUS
CN Acetamide, N-[3-[3-[[[[(tetrahydro-2-furanyl)methyl]amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

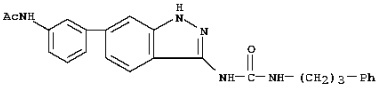


RN 627858-30-4 CAPLUS
CN Acetamide, N-[3-[3-[[[(2-phenylethyl)amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

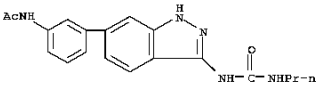
L7 ANSWER 39 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)



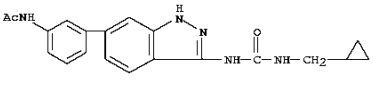
RN 627858-31-5 CAPLUS
CN Acetamide, N-[3-[3-[[[(3-phenylpropyl)amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)



RN 627858-32-6 CAPLUS
CN Acetamide, N-[3-[3-[[(propylamino)carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

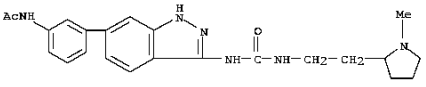


RN 627858-33-7 CAPLUS
CN Acetamide, N-[3-[3-[[[(cyclopropylmethyl)amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

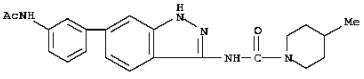


RN 627858-34-8 CAPLUS
CN Acetamide, N-[3-[3-[[[2-(1-methyl-2-pyrrolidinyl)ethyl]amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

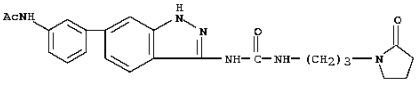
L7 ANSWER 39 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)



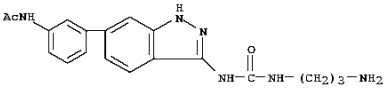
RN 627859-06-7 CAPLUS
CN 1-Piperidinecarboxamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]-4-methyl- (CA INDEX NAME)



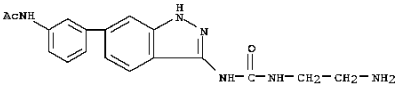
RN 627859-07-8 CAPLUS
CN Acetamide, N-[3-[3-[[[[3-(2-oxo-1-pyrrolidinyl)propyl]amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)



RN 627859-08-9 CAPLUS
CN Acetamide, N-[3-[3-[[[(3-aminopropyl)amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

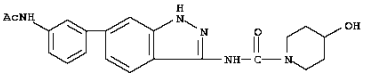


RN 627859-09-0 CAPLUS
CN Acetamide, N-[3-[3-[[[(2-aminoethyl)amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

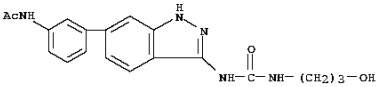


RN 627859-10-3 CAPLUS
CN 1-Piperidinecarboxamide, N-[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]-4-

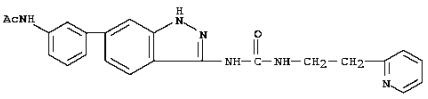
L7 ANSWER 39 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)
hydroxy- (CA INDEX NAME)



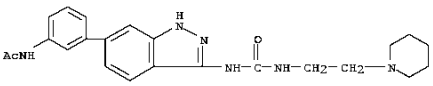
RN 627859-11-4 CAPLUS
CN Acetamide, N-[3-[3-[[[(3-hydroxypropyl)amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)



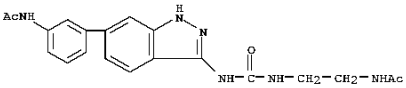
RN 627859-12-5 CAPLUS
CN Acetamide, N-[3-[3-[[[[2-(2-pyridinyl)ethyl]amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)



RN 627859-13-6 CAPLUS
CN Acetamide, N-[3-[3-[[[[2-(1-piperidinyl)ethyl]amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

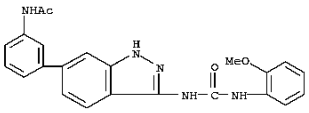


RN 627859-14-7 CAPLUS
CN Acetamide, N-[3-[3-[[[[2-(acetylamino)ethyl]amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)

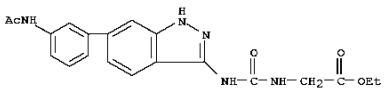


L7 ANSWER 39 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)

IT 627859-44-3P 627859-45-4P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of 3-aminoindazole derivs. as kinase inhibitors)
RN 627859-44-3 CAPLUS
CN Acetamide, N-[3-[3-[[[(2-methoxyphenyl)amino]carbonyl]amino]-1H-indazol-6-yl]phenyl]- (CA INDEX NAME)



RN 627859-45-4 CAPLUS
CN Glycine, N-[[[6-[3-(acetylamino)phenyl]-1H-indazol-3-yl]amino]carbonyl]-, ethyl ester (CA INDEX NAME)



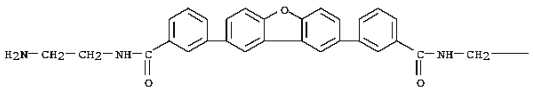
REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 40 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2003:930976 CAPLUS
DOCUMENT NUMBER: 139:395808
TITLE: Preparation of indolylcarbazoles and related compounds as antibacterials and antifungals.
INVENTOR(S): Roberts, Christoher Don; Keicher, Jesse Daniel; Gazginci, Mikail Hakan; Velligan, Mark Douglas
PATENT ASSIGNEE(S): Genelabs Technologies, Inc., USA
SOURCE: U.S. Pat. Appl. Publ., 23 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003220340	A1	20031127	US 2003-438377	20030515
US 6951880	B2	20051004		
WO 2004047724	A2	20040610	WO 2003-US15415	20030515
WO 2004047724	A3	20040729		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003302218	A1	20040618	AU 2003-302218	20030515
PRIORITY APPLN. INFO.:			US 2002-381941P P	20020516
			WO 2003-US15415 W	20030515

OTHER SOURCE(S): MARPAT 139:395808
IT 627075-39-2P 627075-42-7P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of indolylcarbazoles and related compds. as antibacterials and antifungals)
RN 627075-39-2 CAPLUS
CN Benzamide, 3,3'-(2,8-dibenzofurandiyl)bis[N-(2-aminoethyl)- (CA INDEX NAME)

PAGE 1-A



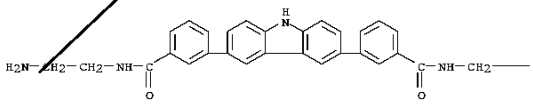
L7 ANSWER 40 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)

PAGE 1-B

—CH2—NH2

RN 627075-42-7 CAPLUS
CN Benzamide, 3,3'-(9H-carbazole-3,6-diyl)bis[N-(2-aminoethyl)- (CA INDEX NAME)

PAGE 1-A



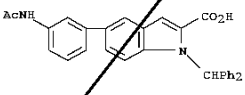
PAGE 1-B

—CH2—NH2

L7 ANSWER 41 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2003:784629 CAPLUS
DOCUMENT NUMBER: 139:292147
TITLE: Preparation of indole derivatives as phospholipase enzyme inhibitors
INVENTOR(S): Seehra, Jasbir S.; Kaila, Neelu; McKew, John C.; Bemis, Jean E.; Xiang, Yibin; Chen, Lihren
PATENT ASSIGNEE(S): Genetics Institute LLC, USA
SOURCE: U.S., 81 pp., Cont.-in-part of U.S. Ser. No. 30,102.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6630496	B1	20031007	US 2000-645042	20000824
BR 9909242	A	20001114	BR 1999-9242	19990217
PRIORITY APPLN. INFO.:			US 1997-918400 B2	19970826
			US 1998-30102 B2	19980225
			WO 1999-IS3388 W	19990217

OTHER SOURCE(S): MARPAT 139:292147
IT 241489-72-5P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of indole derivs. as phospholipase enzyme inhibitors for treatment of inflammatory conditions)
RN 241489-72-5 CAPLUS
CN 1H-Indole-2-carboxylic acid, 5-[3-(acetylamino)phenyl]-1-(diphenylmethyl)- (CA INDEX NAME)



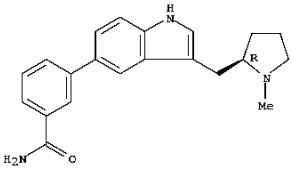
REFERENCE COUNT: 70 THERE ARE 70 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 59 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1994:217271 CAPLUS
DOCUMENT NUMBER: 120:217271
TITLE: Indole derivatives as 5-HT1 agonists
INVENTOR(S): Brown, Alan Daniel; Dickinson, Roger Peter; Wythes, Martin James
PATENT ASSIGNEE(S): Pfizer Ltd., UK; Pfizer Inc.; Pfizer Research and Development Co., N.V./S.A.
SOURCE: PCT Int. Appl., 146 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9321178	A1	19931028	WO 1993-EP867	19930408
W: AU, BR, CA, CZ, FI, HU, JP, KR, NO, NZ, PL, RU, SK, UA, US				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
AU 9339526	A	19931118	AU 1993-39526	19930408
EP 636129	A1	19950201	EP 1993-908928	19930408
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
JP 07502537	T	19950316	JP 1993-517960	19930408
JP 2540017	B2	19961002		
ZA 9302582	A	19941013	ZA 1993-2582	19930413
CN 1082040	A	19940216	CN 1993-105704	19930414
US 5502065	A	19960326	US 1994-307566	19940921
FI 9404805	A	19941012	FI 1994-4805	19941012
PRIORITY APPLN. INFO.:			GB 1992-8161 A	19920414
			WO 1993-EP867 A	19930408

OTHER SOURCE(S): MARPAT 120:217271
IT 153434-71-0 153434-73-2 153434-94-7
153434-99-2 153435-02-0 153435-04-2
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation as 5-HT1 receptor agonist)
RN 153434-71-0 CAPLUS
CN Benzamide, 3-[3-[(1-methyl-2-pyrrolidinyl)methyl]-1H-indol-5-yl]-, (R)- (9CI) (CA INDEX NAME)

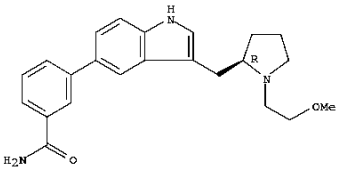
Absolute stereochemistry.



RN 153434-73-2 CAPLUS
CN Benzamide, N-methyl-3-[3-[(1-methyl-2-pyrrolidinyl)methyl]-1H-indol-5-yl]-, (R)- (9CI) (CA INDEX NAME)

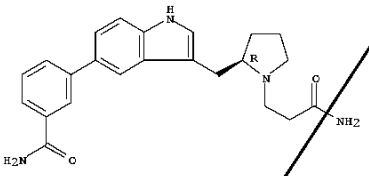
L7 ANSWER 59 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)
CN Benzamide, 3-[3-[[1-(2-methoxyethyl)-2-pyrrolidinyl]methyl]-1H-indol-5-yl]-, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



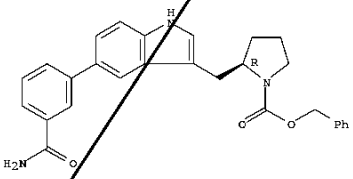
RN 153435-04-2 CAPLUS
CN 1-Pyrrolidinepropanamide, 2-[[5-[3-(aminocarbonyl)phenyl]-1H-indol-3-yl]methyl]-, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



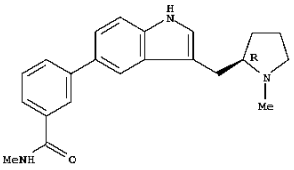
IT 153435-57-5
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation as intermediate in preparation of 5-HT1 receptor agonists)
RN 153435-57-5 CAPLUS
CN 1-Pyrrolidinecarboxylic acid, 2-[[5-[3-(aminocarbonyl)phenyl]-1H-indol-3-yl]methyl]-, phenylmethyl ester, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



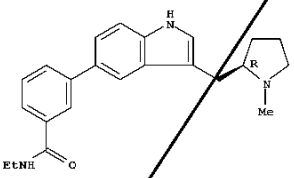
L7 ANSWER 59 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)

Absolute stereochemistry.



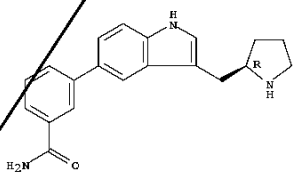
RN 153434-94-7 CAPLUS
CN Benzamide, N-ethyl-3-[3-[(1-methyl-2-pyrrolidinyl)methyl]-1H-indol-5-yl]-, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 153434-99-2 CAPLUS
CN Benzamide, 3-[3-(2-pyrrolidinylmethyl)-1H-indol-5-yl]-, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 153435-02-0 CAPLUS

L7 ANSWER 59 OF 59 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)